

**WEST**[Help](#)[Logout](#)[Interrupt](#)[Main Menu](#)[Search Form](#)[Posting Counts](#)[Show S Numbers](#)[Edit S Numbers](#)[Preferences](#)[Cases](#)**Search Results -**

Terms	Documents
L10 and nitromethyl	2

**Database:**

US Patents Full-Text Database  
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 JPO Abstracts Database  
 EPO Abstracts Database  
 Derwent World Patents Index  
 IBM Technical Disclosure Bulletins

**Search:**

L11

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**DATE:** Friday, December 06, 2002    [Printable Copy](#)    [Create Case](#)

**Set Name**   **Query**  
 side by side

**Hit Count**   **Set Name**  
 result set

DB=USPT,PGPB,JPAB,EPAB,DWPI,TDBD; PLUR=YES; OP=ADJ

<u>L11</u>	L10 and nitromethyl	2	<u>L11</u>
<u>L10</u>	L7 and cyclohexaneacetic acid	17	<u>L10</u>
<u>L9</u>	L8 and nitromethyl	2	<u>L9</u>
<u>L8</u>	L7 and gabapentin	28	<u>L8</u>
<u>L7</u>	562/507	621	<u>L7</u>
<u>L6</u>	560/507	1	<u>L6</u>
<u>L5</u>	560/507	1	<u>L5</u>
<u>L4</u>	nitromethyladj11cyclohexyladj1acetic acid	0	<u>L4</u>
<u>L3</u>	nitromethyladj1cyclohexyladjacetic acid	0	<u>L3</u>
<u>L2</u>	nitromethyl\$1cyclohexyl\$1acetic acid	0	<u>L2</u>
<u>L1</u>	5091567 and nitromethyl\$1cyclohexyl\$1acetic acid	0	<u>L1</u>

END OF SEARCH HISTORY

**WEST**[Generate Collection](#)[Print](#)**Search Results - Record(s) 1 through 2 of 2 returned.**☐ 1. Document ID: US 6153650 A

L11: Entry 1 of 2

File: USPT

Nov 28, 2000

US-PAT-NO: 6153650

DOCUMENT-IDENTIFIER: US 6153650 A

TITLE: Substituted gamma aminobutyric acids as pharmaceutical agents

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draw Desc	Image
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☐ 2. Document ID: US 5091567 A

L11: Entry 2 of 2

File: USPT

Feb 25, 1992

US-PAT-NO: 5091567

DOCUMENT-IDENTIFIER: US 5091567 A

TITLE: Process for the preparation of 1-aminomethyl-1-cyclohexaneacetic acid

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draw Desc	Image
------	-------	----------	-------	--------	----------------	------	-----------	-----------	-------------	--------	-----	-----------	-------

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Terms	Documents
L10 and nitromethyl	2

**Display Format:**

-

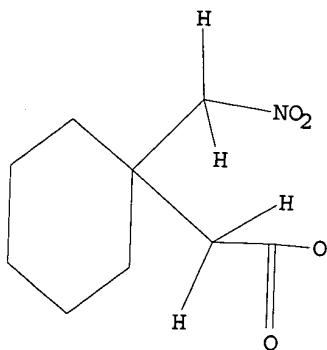
[Change Format](#)[Previous Page](#)[Next Page](#)

L6      STRUCTURE UPLOADED

=> d l6

L6 HAS NO ANSWERS

L6                      STR



Structure attributes must be viewed using STN Express query preparation.

=> s l6

**REGISTRY INITIATED**

Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

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SAMPLE SEARCH INITIATED. 10:46:59 FILE 'REGISTRY'  
SAMPLE SCREEN SEARCH COMPLETED -            5 TO ITERATE

100.0% PROCESSED            5 ITERATIONS  
SEARCH TIME: 00.00.01

0 ANSWERS

FULL FILE PROJECTIONS:    ONLINE    \*\*COMPLETE\*\*  
                             BATCH    \*\*COMPLETE\*\*  
PROJECTED ITERATIONS:            5 TO            234  
PROJECTED ANSWERS:                0 TO            0

L7                      0 SEA SSS SAM L6

L8                      0 L7

=> s l6 full

**REGISTRY INITIATED**

Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

FULL SEARCH INITIATED 10:47:09 FILE 'REGISTRY'  
FULL SCREEN SEARCH COMPLETED -            69 TO ITERATE

100.0% PROCESSED            69 ITERATIONS  
SEARCH TIME: 00.00.01

10 ANSWERS

L9                      10 SEA SSS FUL L6

L10 8 L9

=> s l10 and (benzyl or diphenylmethyl)

137602 BENZYL

3296 DIPHENYLMETHYL

L11 2 L10 AND (BENZYL OR DIPHENYLMETHYL)

=> d 1-2 ibib abs hitstr

L11 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 2000:457016 CAPLUS

DOCUMENT NUMBER: 133:59094

TITLE: Process for the synthesis of 1-(aminomethyl)cyclohexylacetic acid

INVENTOR(S): Gizur, Tibor; Lengyel, Zoltanne; Szalai, Krisztina

PATENT ASSIGNEE(S): Richter Gedeon Vegyeszeti Gyar Rt., Hung.

SOURCE: PCT Int. Appl., 13 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000039074	A1	20000706	WO 1999-HU102	19991223
W:	AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
EP 1140793	A1	20011010	EP 1999-963652	19991223
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, LT, LV, FI, RO			
JP 2002533431	T2	20021008	JP 2000-590987	19991223
PRIORITY APPLN. INFO.:			HU 1998-3034	A 19981229
			WO 1999-HU102	W 19991223

OTHER SOURCE(S): MARPAT 133:59094

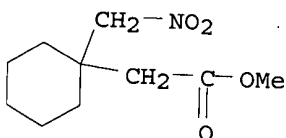
AB 1-(Aminomethyl)cyclohexylacetic acid (gabapentin) was prepd. via hydrogenation of intermediate 1-(nitromethyl)cyclohexylacetic acid (I) or benzyl or diphenylmethyl esters. Thus, a soln. of I in MeOH was hydrogenated in the presence of 0.2 g Pd on activated carbon at atm. pressure to afford 80% gabapentin.

IT 112777-75-0 277333-40-1 277333-41-2  
277333-42-3

RL: RCT (Reactant); RACT (Reactant or reagent)  
(synthesis of aminomethylcyclohexylacetic acid)

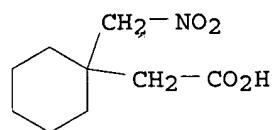
RN 112777-75-0 CAPLUS

CN Cyclohexaneacetic acid, 1-(nitromethyl)-, methyl ester (9CI) (CA INDEX NAME)

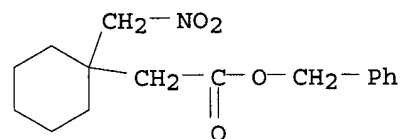


RN 277333-40-1 CAPLUS

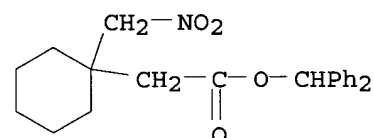
CN Cyclohexaneacetic acid, 1-(nitromethyl)- (9CI) (CA INDEX NAME)



RN 277333-41-2 CAPLUS  
 CN Cyclohexaneacetic acid, 1-(nitromethyl)-, phenylmethyl ester (9CI) (CA  
 INDEX NAME)



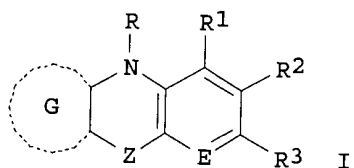
RN 277333-42-3 CAPLUS  
 CN Cyclohexaneacetic acid, 1-(nitromethyl)-, diphenylmethyl ester (9CI) (CA  
 INDEX NAME)



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS  
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2002 ACS  
 ACCESSION NUMBER: 1998:126254 CAPLUS  
 DOCUMENT NUMBER: 128:204878  
 TITLE: Preparation of pyrazinobenzothiazine derivatives and  
 analogs for the treatment of inflammation and  
 autoimmune diseases  
 INVENTOR(S): Kaneko, Toshihiko; Clark, Richard; Ohi, Norihito;  
 Ozaki, Fumihiro; Kawahara, Tetsuya; Kamada, Atsushi;  
 Okano, Kazuo; Yokohama, Hiromitsu; Muramoto, Kenzo;  
 Arai, Tohru; Ohkuro, Masayoshi; Takenaka, Osamu;  
 Sonoda, Jiro  
 PATENT ASSIGNEE(S): Eisai Co., Ltd., Japan; Kaneko, Toshihiko; Clark,  
 Richard; Ohi, Norihito; Ozaki, Fumihiro; Kawahara,  
 Tetsuya; Kamada, Atsushi; Okano, Kazuo; Yokohama,  
 Hiromitsu; et al.  
 SOURCE: PCT Int. Appl., 1344 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9806720	A1	19980219	WO 1997-JP2787	19970808
W: AU, CA, CN, HU, JP, KR, MX, NO, NZ, RU, US				
RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
AU 9737849	A1	19980306	AU 1997-37849	19970808
ZA 9707103	A	19990208	ZA 1997-7103	19970808
EP 934941	A1	19990811	EP 1997-934750	19970808
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, FI				
PRIORITY APPLN. INFO.:			JP 1996-210344	19960809
			WO 1997-JP2787	19970808
OTHER SOURCE(S):		MARPAT 128:204878		
GI				



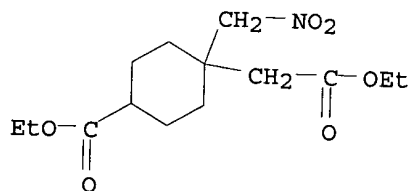
AB The title compds. I [R1 to R3 are the same or different and each represents hydrogen, optionally substituted lower alkyl, optionally substituted cycloalkyl, etc., provided that when R1 to R3 are all optionally substituted lower alkyl groups, they do not simultaneously represent Me groups; R represents hydrogen, lower alkyl, etc.; E represents N, C, etc.; Z represents O, S, SO, SO2, etc.; and the ring G represents an optionally substituted heteroaryl ring having at least one nitrogen atom] are prepd. I are useful in the treatment and prevention of inflammatory immunol. diseases, autoimmune diseases, rheumatism, collagen disease, asthma, nephritis, ischemic reflow disorders, psoriasis, atopic dermatitis or rejection reactions following organ transplantation. The compd. (syn)-[3-(10H-pyrazino[2,3-b][1,4]benzothiazin-8-ylmethyl)-3-azabicyclo[3.3.1]nona-9-yl]acetic acid (II) at 10 mg/kg orally gave 65% inhibition of carrageenin-induced inflammation in rats. II in vitro showed IC50 of 2.3 .mu.M against the expression of ICAM-1.

IT 203661-25-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)  
(prepn. of pyrazinobenzothiazine derivs. and analogs for treatment of inflammation and autoimmune diseases)

RN 203661-25-0 CAPLUS

CN Cyclohexaneacetic acid, 4-(ethoxycarbonyl)-1-(nitromethyl)-, ethyl ester  
(9CI) (CA INDEX NAME)



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